

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
3 October 2002 (03.10.2002)

PCT

(10) International Publication Number
WO 02/076230 A1(51) International Patent Classification⁷: **A23G 3/30**,
C08G 63/08, 63/02**Robson** [US/US]; 111 Holly Dr., Hattiesburg, MS 39402 (US).

(21) International Application Number: PCT/DK02/00203

(74) Agent: **PATENTGRUPPEN APS**; Arosgaarden,
Aaboulevarden 31, DK-8000 Aarhus C (DK).

(22) International Filing Date: 25 March 2002 (25.03.2002)

(25) Filing Language: English

(81) Designated States (*national*): AE, AG, AL, AM, AT, AT (utility model), AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, CZ (utility model), DE, DE (utility model), DK, DK (utility model), DM, DZ, EC, EE, EE (utility model), ES, FI, FI (utility model), GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (utility model), SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW.

(26) Publication Language: English

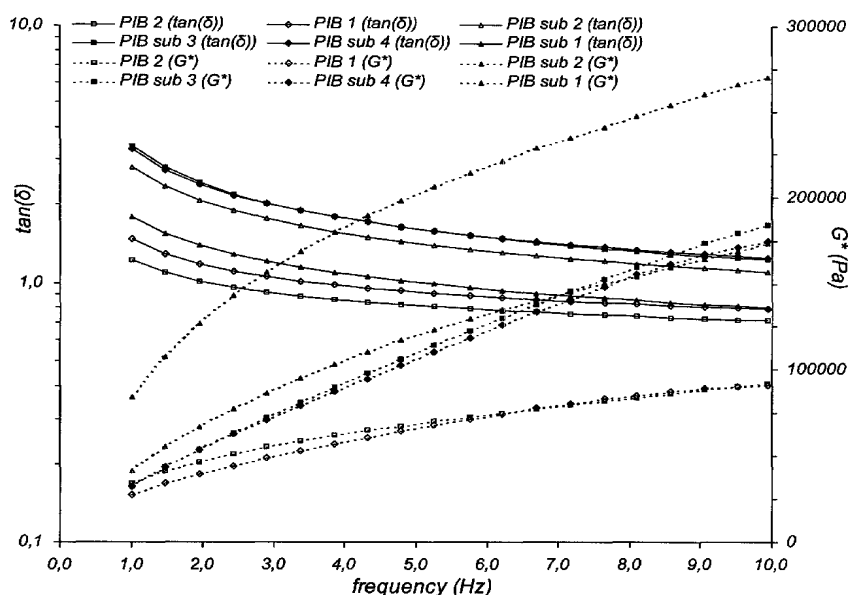
(30) Priority Data:
PA 2001 00494 23 March 2001 (23.03.2001) DK
60/303,098 6 July 2001 (06.07.2001) US(71) Applicant (*for all designated States except US*): **DANDY**
A/S [DK/DK]; Dandyvej 19, DK-7100 Vejle (DK).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): **WITTORFF, Helle** [DK/DK]; Hegnsgårdsvej 63, DK-7120 Vejle (DK). **ANDERSEN, Lone** [DK/DK]; Gl. Strandvej 1, DK-5500 Middelfart (DK). **ISAKSEN, Anette** [DK/DK]; Rønningesminde 22, DK-6000 Kolding (DK). **STOREY,**(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent

[Continued on next page]

(54) Title: DEGRADABLE ELASTOMERS FOR CHEWING GUM BASE

(57) Abstract: There is provided a novel degradable gum base comprising low molecular weight elastomer replacement compounds which are generally applicable for chewing gum formulations. In particular there is provided a gum base and a chewing gum comprising a polyester polymer obtainable by the polymerisation of two or more different cyclic ester monomers, wherein the cyclic ester monomers have a low glass transition temperature (T_g) and the polyester polymer has a glass transition temperature (T_g) in the range from (-20 °C) to (-80 °C).

WO 02/076230 A1



(BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

Published:

- *with international search report*
- *before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments*

DEGRADABLE ELASTOMERS FOR CHEWING GUM BASE

FIELD OF THE INVENTION

5 The present invention pertains to the field of chewing gum. In particular, there is provided a novel degradable gum base comprising low molecular weight elastomer replacement compounds which are generally applicable for chewing gum formulations. In particular the present invention provides a gum base and a chewing gum comprising a polyester polymer obtainable by the polymerisation of two or
10 more different cyclic ester monomers, wherein the cyclic ester monomers have a low glass transition temperature (T_g) and the polyester polymer has a glass transition temperature (T_g) in the range from (-20°C) to (-80°C) .

TECHNICAL BACKGROUND AND PRIOR ART

15 It is generally recognized that chewing gum that is dropped in indoor or outdoor environments gives rise to considerable nuisances and inconveniences due to fact that the dropped gum sticks firmly to e.g. street and pavement surfaces and to shoes and clothes of people being present or moving in the environments. Adding
20 substantially to such nuisances and inconveniences is the fact that currently available chewing gum products are based on the use of elastomeric and resinous polymers of natural or synthetic origin that are substantially non-degradable in the environment.

City authorities and others being responsible for cleanliness of indoor and outdoor
25 environments therefore have to exercise considerable efforts to remove dropped chewing gum, such efforts, however, being both costly and without satisfactory results.

There have been attempts to reduce the nuisances associated with the widespread use
30 of chewing gum e.g. by improving cleaning methods to make them more effective with regard to removal of dropped chewing gum remnants or by incorporating anti-

sticking agents into chewing gum formulations. However, none of these precautions have contributed significantly to solving the pollution problem.

The past two decades have seen an increasing amount of interest paid to synthetic polyesters for a variety of applications ranging from biomedical devices to gum bases. Many of these polymers are degradable and readily hydrolyse to their monomeric hydroxy-acids, which are easily removed by metabolic pathways. Degradable (also referred to as biodegradable) polymers are e.g. anticipated as alternatives to traditional non- or low-degradable plastics such as poly(styrene), poly(isobutylene), and poly(methyl-methacrylate).

Thus, it has recently been disclosed, e.g. in US 5,672,367 that chewing gum may be made from certain synthetic polymers having in their polymer chains chemically unstable bonds that can be broken under the influence of light or hydrolytically into water-soluble and non-toxic components. The claimed chewing gum comprises at least one degradable polyester polymer obtained by the polymerisation of cyclic esters, e.g. based on lactides, glycolides, trimethylene carbonate and ϵ -caprolactone. It is mentioned in this patent that chewing gum made from such polymers that are referred to as biodegradable are degradable in the environment.

US 6,153,231 discloses degradable chewing gum bases which comprises poly(lactic acid) co-polymers selected from poly(lactid acid-dimer-fatty acid-oxazoline) copolymers and poly(lactic acid-diol-urethane) copolymers.

In general, a chewing gum composition typically comprises a water-soluble bulk portion, a water-insoluble gum base portion and typically water-insoluble flavouring agents.

The water-insoluble gum base generally comprises one or more elastomeric compounds which may be of synthetic or natural origin, one or more resinous compounds, one or more elastomer plasticizers, fillers, softening compounds and

minor amounts of miscellaneous ingredients such as antioxidants and colorants and others.

Elastomers provide the rubbery, cohesive nature to the gum base which varies
5 depending on this components chemical structure and how it is blended with other ingredients. Typically, the elastomeric compounds in gum base are non-degradable. Such elastomers includes synthetic elastomers such as polyisobutylene, isobutylene-isoprene copolymer (butyl elastomer), styrene-butadiene copolymers, polyisoprene, polyethylene, polyvinylacetate, vinyl acetate-vinyl laurate copolymer and combinations hereof. However, also natural elastomers are presently applied in chewing
10 gum bases. Such natural elastomers may include natural rubber such as smoked or liquid latex and guayule, natural gums such as jelutong, lechi caspi perillo, massaranduba balata, massaranduba chocolate, nispero, rosidinha, chicle, gutta percha, gutta kataiu, niger gutta, tunu, chilte, chiquibul and gutta hang kang.

15 It has now been found by the present inventors, that it is possible, in a chewing gum base, to replace the elastomeric compounds, such as e.g. polyisobutylene, with a degradable polymer comprising a polyester polymer obtainable by the polymerisation of two or more different cyclic ester monomers, wherein the cyclic
20 ester monomers have a low glass transition temperature (T_g) and the polyester polymer has a glass transition temperature (T_g) in the range from (-20°C) to (-80°C). Thus, it has surprisingly been found that chewing gum bases prepared with such degradable polymers have the same or similar rheological properties (such as plasticity (storage modulus) and elasticity (loss modulus)) as e.g. conventional gum
25 bases prepared with polyisobutylene (PIB).

As elastomeric compounds typically constitutes between 20 to 60% of the entire gum base composition, the replacement of this component of the gum base with a degradable component highly improve the general degradability of the gum base and
30 hence the chewing gum as such.

SUMMARY OF THE INVENTION

Accordingly, an aspect of the present invention pertains in a gum base comprising a polyester polymer obtainable by the polymerisation of two or more different cyclic ester monomers, wherein the cyclic ester monomers have a low glass transition temperature (T_g) and the polyester polymer has a glass transition temperature (T_g) in the range from (-20°C) to (-80°C).

10 In a still further aspect the invention relates to a chewing gum comprising a chewing gum base as defined above and below herein.

According to a further embodiment of the invention, a chewing gum or a gum base may comprise a partly substituted functional group, here an elastomer and where the substituted functional group is bio-degradable.

According to a further embodiment of the invention, it has been determined that conventional non-biodegradable functional groups as such may be substituted by other rheologically matching bio-degradable polymers.

DETAILED DISCLOSURE

A strategy for creation of an elastomer for a gum base is to create a polymer that has a low glass transition temperature and is either totally amorphous or is slightly
5 crystalline with a crystalline melting temperature below room temperature.

A preferred way, to obtain such a polymer, is to use two or more low-Tg monomers, in combination, so that the dissimilar repeating units hinder crystallization.

10 Accordingly, an aspect of the invention relates to a gum base comprising a polyester polymer obtainable by the polymerisation of two or more different cyclic ester monomers, wherein the cyclic ester monomers have a low glass transition temperature (Tg) and the polyester polymer has a glass transition temperature (Tg) in the range from (-20°C) to (-80°C).

15 Preferably, the cyclic ester monomers are selected from the group consisting of a 4-membered lactone, a 5-membered lactone, a 6-membered lactone, a 7-membered lactone, a 8-membered lactone, a 5-membered cyclic carbonate and a 6-membered cyclic carbonate.

20 The lactone is preferably selected from the group consisting of β -propiolactone, γ -butyrolactone, δ -valerolactone, ϵ -caprolactone and 7-heptanolactone; and the cyclic carbonate is preferably an ethylene carbonate or a trimethylene carbonate.

25 A preferred embodiment relates to a gum base comprising a polyester polymer obtainable by the polymerisation of two or more different cyclic ester monomers, wherein the cyclic ester monomers are selected from the group consisting of ϵ -caprolactone, δ -valerolactone and trimethylene carbonate.

30 The cyclic ester monomer ϵ -caprolactone is a preferred monomer and preferably the polyester polymer contain at least 50 mole% of ϵ -caprolactone.

Further, the polyester polymer has preferably a glass transition temperature (T_g) in the range from (-25°C) to (-75°C), more preferably the polyester polymer has a glass transition temperature (T_g) in the range from (-45°C) to (-75°C).

- 5 Below is described preferred embodiments relating to a gum base comprising a poly(ε-caprolactone-co-δ-valerolactone) and a gum base comprising a poly(ε-caprolactone-co-δ-valerolactone-co-trimethylene carbonate).

The different embodiments of a gum base below are also relevant for a gum base as described above.

10

It is a major objective of the present invention to provide gum bases for chewing gum which results in chewing gum products that following chewing are more readily degraded in the environment if improperly dropped or discarded here by the user and/or which, relative to chewing gum comprising conventional non-degradable
15 polymers can be removed more readily mechanically and/or by the use of cleaning agents.

20

Accordingly, the chewing gum base provided herein is a gum base which when applied in chewing gum, renders the chewing gum more capable of undergoing a physical, chemical and/or biological degradation whereby e.g. dumped chewing gum waste becomes more readily removable from the site of dumping or is eventually disintegrated to lumps or particles which are no longer recognisable as being chewing gum remnants. The degradation or disintegration of the gum base provided herein can be effected or induced by physical factors such as temperature, light,
25 moisture, by chemical factors such as hydrolysis caused by a change in pH or by the action of appropriate enzymes capable of degrading the co-polymers according to the invention.

30

Accordingly, it is one objective of the present invention to provide a gum base comprising a degradable co-polymer consisting of ε-caprolactone and δ-valerolactone and a terpolymer consisting of ε-caprolactone, δ-valerolactone and trimethylene carbonate.

As mentioned above, it has been found possible, by applying such a co-polymer or terpolymer, to completely replace a synthetic and substantially non-degradable elastomeric compound such as polyisobutylene (PIB), which is typically applied in chewing gum compositions. Surprisingly, as will appear from the following examples, that by matching the rheological profil of the polyisobutylene with the rheological profil of a degradable co- or ter-polymers of ϵ -caprolactone, δ -valerolactone or trimethylene carbonate or mixtures thereof, then this replacement can be made without impairing the rheological properties of the gum base and the chewing gum made from such gum bases. Thus, it is possible to obtain rheological properties (such as plasticity (loss modulus) and elasticity (storage modulus)), which are similar to conventional gum bases prepared with PIB. Plasticity and elasticity are parameters that are essential for the texture in the final chewing gum.

It is contemplated that the above novel degradable polymers advantageously may be applied as elastomer replacement for other elastomeric compounds than polyisobutylene. Accordingly, it is also within the scope of the invention that the polymers poly(ϵ -caprolactone-co- δ -valerolactone) and poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) may be applied as replacements for elastomeric compounds such as isobutylene-isoprene copolymer (butyl elastomer), styrene-butadiene copolymers, polyisoprene, polyethylene, polyvinyleacetate, vinyl acatate vinyl laurate copolymer and combinations thereof.

Thus, it is one objective of the present invention to provide a chewing gum base comprising poly(ϵ -caprolactone-co- δ -valerolactone) co-polymer.

The preparation of the poly(ϵ -caprolactone-co- δ -valerolactone) co-polymer may be performed by various suitable polymerisation processes which are well known in the art, e.g. by ring opening polymerisation (ROP) in the presence of an appropriate catalyst. Accordingly, in one embodiment stannous octoate (SO) may advantageously be applied as a catalyst and a low molecular weight alcohol (e.g. propylene glycol) as initiator to polymerise a mixture of ϵ -caprolactone and δ -

valerolactone monomers and in order to obtain poly(ϵ -caprolactone-co- δ -valerolactone). However, it is also contemplated that the polymerisations may be mediated by applying various aluminum-alkoxide compounds as initiators.

- 5 It will be appreciated that the mol percentage of the monomers in the polymers of the present invention may be individually adjusted, by applying different polymerisation conditions, in order to obtain the desired rheological characteristics of the gum base in which the polymer is intended to be applied. Thus, it is contemplated that a wide range of mol percentages of the individual monomers may be advantageously
10 applied.

Accordingly, in a useful embodiment, the poly(ϵ -caprolactone-co- δ -valerolactone) co-polymer may be synthesised to have a specific mol percentage of each of the monomers. Thus, in one embodiment of the invention, the mol percentage of ϵ -
15 caprolactone in the poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 1 - 99 mol %. The mol percentage of the individual monomers of the synthesised polymers may e.g. be determined by means of e.g. ^{13}C NMR-analysis.

In a further embodiment the mol percentage of ϵ -caprolactone in the poly(ϵ -
20 caprolactone-co- δ -valerolactone) is in the range of 40 - 80 mol %, including the range of 50 - 70 mol %, such as the range of 55 - 65 mol %. In one embodiment, the mol percentage of ϵ -caprolactone in the poly(ϵ -caprolactone-co- δ -valerolactone) is about 60 mol %.

25 Likewise, it will be appreciated that the chewing gum base according to the invention advantageously may comprise poly(ϵ -caprolactone-co- δ -valerolactone) wherein the mol percentage of δ -valerolactone is in the range of 1 - 99 mol %, including the range of 20 - 60 mol %, such as the range of 30 - 50 mol %. In one embodiment the mol percentage of δ -valerolactone is about 40 mol %.

30

As mentioned above, it is contemplated that a suitable gum base may comprise poly(ϵ -caprolactone-co- δ -valerolactone) having different structural characteristics

such as molecular weight including number average molecular weight (M_n) and weight average molecular weight (M_w). Accordingly, in one embodiment the chewing gum base according to the invention comprises poly(ϵ -caprolactone-co- δ -valerolactone) having a number average molecular weight (M_n) in the range of 10,000 - 125,000 g/mol, including the range of 20,000 - 100,000 g/mol, such as the range of 30,000 - 90,000 g/mol, including the range of 40,000 - 80,000 g/mol.

An important rheological feature for gum bases which are applied in chewing gum compositions, is the glass transition temperature (T_g). As used herein, the glass transition temperature means the temperature at which the ratio of the storage modulus G' (elasticity) and the loss modulus G'' (plasticity) equals one. Storage modulus G' and loss modulus G'' of polymers may in general be determined by applying a rheometer such as AR1000 from AT Instruments.

In one embodiment the gum base according to the invention comprises a poly(ϵ -caprolactone-co- δ -valerolactone) co-polymer having a glass transition temperature (T_g) which is less than 0°C. In useful embodiments, the glass transition temperature of the poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of (-40°C) - (-80°C), including the range of (-50°C) - (-70°C).

The crystallinity may be depressed by incorporating chain branching or introducing a co-monomer. Possible biodegradable co-monomers include δ -valerolactone (VAL), a six-membered cyclic ester, and trimethylene carbonate (TMC), a six-membered cyclic carbonate

As mentioned above, it is a further objective of the present invention to provide a chewing gum base comprising poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate). Accordingly, in a further aspect the degradable co-monomer trimethylene carbonate is included in the polymer according to the invention.

The preparation of poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) terpolymer may e.g. be performed by the above mentioned various polymerisation processes which are well known in the art.

- 5 In accordance with the invention, the mol percentage of ϵ -caprolactone in the poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) may in useful embodiments be in the range of 1 - 99 mol %, such as the range of 20 - 80 mol %, including the range of 40 - 60 mol %. In a presently preferred embodiment the mol percentage of ϵ -caprolactone in the poly(ϵ -caprolactone-co- δ -valerolactone-co-
10 trimethylene carbonate) is about 50 mol %.

Also in accordance with the invention, the gum base may comprise poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) having a mol percentage of δ -valerolactone in the range of 1 - 99 mol %, including the range of 20 - 60 mol
15 %, such as in the range of 30 - 50 mol %. In one specific embodiment, the mol percentage of δ -valerolactone in the poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is about 40 mol %.

The mol percentage of trimethylene carbonate in the poly(ϵ -caprolactone-co- δ -
20 valerolactone-co-trimethylene carbonate) may in useful embodiments be in the range of range of 1 - 50 mol %, including the range of 2 - 30 mol %, such as the range of 5 - 15 mol %. In a useful embodiment, the mol percentage of trimethylene carbonate in the poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is about 10 mol %.

25

As mentioned above, a structural characteristic such as molecular weight may be tailored for each specific gum base. Accordingly, in one embodiment the chewing gum base comprises a poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) with a average number molecular weight (M_n) in the range of 10,000 -
30 150,000 g/mol. In useful embodiments the molecular weight (M_n) of the poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 20,000

- 100,000 g/mol, including the range of 30,000 - 90,000 g/mol, such as the range of 40,000 - 80,000 g/mol.

In further useful embodiments, the gum base of the invention comprises poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) with a glass transition temperature T_g of less than 0°C. However, it is also within the scope of the invention that the glass transition temperature T_g of the poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of (-40°C) - (-80°C), including the range of (-50°C) - (-75°C).

10

As was mentioned above, the present invention also provides a chewing gum comprising a chewing gum base comprising poly(ϵ -caprolactone-co- δ -valerolactone) co-polymer or comprising poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) terpolymer. However, it will be appreciated that in specific embodiments the co-polymer and the terpolymer may advantageously be combined in a gum base in order to achieve specific rheological features or characteristics. Accordingly, there is provided a chewing gum product which is based on the gum base according to the invention which is disclosed herein.

As used herein, the expressions "gum base" refers in general to the water insoluble part of the chewing gum which typically constitutes 10 to 99% by weight (preferably 10 to 50% by weight) of the total chewing gum formulation. Chewing gum base formulations typically comprises one or more elastomeric compounds which may be of synthetic or natural origin, one or more resinous compounds of natural or synthetic origin, fillers, softening compounds and minor amounts of miscellaneous ingredients such as antioxidants and colorants, etc.

Thus, it is within the scope of the invention that the gum base part, in addition to the degradable elastomers co-polymer poly(caprolactone-co- δ -valerolactone) and terpolymer poly(caprolactone-co- δ -valerolactone-co-trimethylene carbonate), contains a proportion of non-degradable polymeric elastomers and/or resins which may be of natural or synthetic origin. The proportion of such non-degradable

30

polymers may be in the range of 1-99% by weight including the range of 5 to 90% by weight such as in the range of 10-50% by weight.

In this context, useful synthetic elastomers include, but are not limited to, synthetic
5 elastomers listed in Food and Drug Administration, CFR, Title 21, Section 172,615, the Masticatory Substances, Synthetic) such as polyisobutylene with a gas pressure chromatography (GPC) average molecular weight in the range of about 10,000 to about 1,000,000 including the range of 50,000 to 80,000, isobutylene-isoprene copolymer (butyl elastomer), styrene-butadiene copolymers e.g. having styrene-
10 butadiene ratios of about 1:3 to about 3:1, polyisoprene, polyethylene, polyvinyl acetat, vinyl acetate-vinyl laurate copolymer e.g. having a vinyl laurate content of about 5 to about 50% by weight such as 10 to 45% by weight of the copolymer, and combinations hereof.

15 It is e.g. common in the industry to combine in a gum base a synthetic elastomer having a high molecular weight and a low-molecular-weight elastomer. Presently preferred combinations of synthetic elastomers include, but are not limited to, polyisobutylene and styrene-butadiene, polyisobutylene and polyisoprene, polyisobutylene and isobutylene-isoprene copolymer (butyl rubber) and a
20 combination of polyisobutylene, styrene-butadiene copolymer and isobutylene isoprene copolymer, and all of the above individual synthetic polymers in admixture with polyvinyl acetate, vinyl acetate-vinyl laurate copolymers, respectively and mixtures thereof.

25 Useful natural non-degradable elastomers include the elastomers listed in Food and Drug Administration, CFR, Title 21, Section 172,615, as "Masticatory Substances of Natural Vegetable Origin" including natural rubber compounds such as smoked or liquid latex and guayule and other natural gums including jelutong, lechi caspi, massaranduba balata, sorva, perillo, rosindinha, massaranduba chocolate, chicle,
30 nispero, gutta hang kang, and combinations thereof. The preferred synthetic elastomer and natural elastomer concentrations vary depending on whether the chewing gum in which the base is used is adhesive or conventional, bubble gum or

regular gum, as discussed below. Presently preferred natural elastomers include jelutong, chicle, massaranduba balata and sorva.

However, it is also contemplated that in useful embodiments, the gum base according to the invention which comprise poly(ϵ -caprolactone-co- δ -valerolactone) and/or poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate), may advantageously further comprise elastomeric or resinous polymer compounds which are environmentally or biologically degradable.

10 In the present context the terms environmentally or biologically degradable polymer compounds refers to chewing gum base components which, after dumping the chewing gum, is capable of undergoing a physical, chemical and/or biological degradation whereby the dumped chewing gum waste becomes more readily removable from the site of dumping or is eventually disintegrated to lumps or
15 particles which are no longer recognisable as being chewing gum remnants. The degradation or disintegration of such degradable polymers can be effected or induced by physical factors such as temperature, light, moisture, by chemical factors such as hydrolysis caused by a change in pH or by the action of enzymes capable of degrading the polymers. In other useful embodiments all of the polymer components
20 of the gum base are environmentally degradable or biodegradable polymers.

Accordingly, suitable examples of additional environmentally or biologically degradable chewing gum base polymers which can be applied in accordance with the gum base of the present invention include degradable polyesters, polycarbonates, polyester amides, polypeptides, homopolymers of amino acids such as polylysine, and
25 proteins including derivatives hereof such as e.g. protein hydrolysates including a zein hydrolysate. Particularly useful compounds of this type include polyester polymers obtained by the polymerisation of one or more cyclic esters such as lactide, glycolide, trimethylene carbonate, δ -valerolactone, β -propiolactone and ϵ -
30 caprolactone. Such degradable polymers may be homopolymers or copolymers, including block-polymers.

In accordance with the invention, the chewing gum base components which are useful may include one or more resinous compounds contributing to obtain the desired masticatory properties and acting as plasticizers for the elastomers of the gum base composition. In the present context, useful elastomer plasticizers include
5 synthetic resins such as polyvinyl acetate (PVAc) having a GPC average molecular weight in the range of 2,000 to about 90,000 such as the range of 3,000 to 80,000, and natural resins such as natural rosin esters, often referred to as ester gums including as examples glycerol esters of partially hydrogenated rosins, glycerol esters of polymerised rosins, glycerol esters of partially dimerised rosins, glycerol esters of
10 tall oil rosins, pentaerythritol esters of partially hydrogenated rosins, methyl esters of rosins, partially hydrogenated methyl esters of rosins, pentaerythritol esters of rosins. Other useful resinous compounds include synthetic resins such as terpene resins derived from alpha-pinene, beta-pinene, and/or d-limonene, natural terpene resins; and any suitable combinations of the foregoing. The preferred elastomer
15 plasticizers will also vary depending on the specific application, and on the type of elastomer(s) being used.

A chewing gum base formulation may, if desired, include one or more fillers/texturisers including as examples, magnesium and calcium carbonate, sodium
20 sulphate, ground limestone, silicate compounds such as magnesium and aluminium silicate, kaolin and clay, aluminium oxide, silicon oxide, talc, titanium oxide, mono-, di- and tri-calcium phosphates, cellulose polymers, such as wood, and combinations thereof.

25 The fillers/texturisers may also include natural organic fibres such as fruit vegetable fibres, grain, rice, cellulose and combinations thereof.

As used herein the term "softener" designates an ingredient, which softens the gum base or chewing gum formulation and encompasses waxes, fats, oils, emulsifiers,
30 surfactants and solubilisers.

A gum base formulation may, in accordance with the present invention comprise one or more fats e.g. tallow, hydrogenated tallow, any completely or partially hydrogenated animal fats, completely hydrogenated and partially hydrogenated vegetable oils or fats, cocoa butter, degreased cocoa butter, glycerol monostearate, glycerol triacetate, lecithin, mono-, di- and triglycerides, acetylated monoglycerides, fatty acids (e.g. stearic, palmitic, oleic and linoleic acids), and/or combinations thereof.

To soften the gum base further and to provide it with water binding properties, which confer to the gum base a pleasant smooth surface and reduce its adhesive properties, one or more emulsifiers is/are usually added to the composition, typically in an amount of 0 to 18% by weight, preferably 0 to 12% weight of the gum base. Mono- and diglycerides of edible fatty acids, lactic acid esters and acetic acid esters of mono- and di- and triglycerides of edible fatty acids, acetylated mono and diglycerides, sucrose polyesters or sugar esters of edible fatty acids including those disclosed in WO 00/25598, which is incorporated herein by reference, Na-, K-, Mg- and Ca-stearates, lecithin, hydroxylated lecithin, glycerol monostearate, glycerol triacetate, fatty acids (e.g. stearic, palmitic, oleic and linoleic acids), propylgallates and combinations thereof are examples of conventionally used emulsifiers which can be added to the chewing gum base. In case of the presence of a biologically or pharmaceutically active ingredient as defined below, the formulation may comprise certain specific emulsifiers and/or solubilisers in order to disperse and release the active ingredient.

Waxes are conventionally used for the adjustment of the consistency and for softening of the chewing gum base when preparing chewing gum bases. In connection with the present invention any conventionally used and suitable type of wax may be used, such as for instance rice bran wax, polyethylene wax, petroleum wax (refined paraffin and microcrystalline wax), paraffin, bees' wax, carnauba wax, and candelilla wax.

Furthermore, the gum base formulation may, in accordance with the present invention, comprise colourants and whiteners such as FD&C-type dyes and lakes, fruit and vegetable extracts, titanium dioxide and combinations thereof. Further useful chewing gum base components include antioxidants, e.g. butylated
5 hydroxytoluene (BHT), butyl hydroxyanisol (BHA), propylgallate and tocopherols, and preservatives.

The composition of chewing gum base formulations which are admixed with chewing gum additives as defined below can vary substantially depending on the
10 particular product to be prepared and on the desired masticatory and other sensory characteristics of the final product. However, typical ranges (weight%) of the above gum base components are: 5 to 100% by weight (e.g. 5 to 50% by weight) elastomeric compounds, 5 to 55% by weight elastomer plasticizers, 0 to 50% by weight filler/texturiser, 5 to 35% by weight softener and 0 to 1% by weight of
15 miscellaneous ingredients such as antioxidants, colorants, etc.

A chewing gum centre formulation comprises, in addition to the above water-insoluble gum base components, a generally water soluble part comprising a range of chewing gum additives. In the present context, the term "chewing gum additive" is
20 used to designate any component, which in a conventional chewing gum manufacturing process is added to the gum base. The major proportion of such conventionally used additives are water soluble, but water-insoluble components, such as e.g. water-insoluble flavouring compounds, can also be included.

25 In the present context, chewing gum additives include bulk sweeteners, high intensity sweeteners, flavouring agents, softeners, emulsifiers, colouring agents, binding agents, acidulants, fillers, antioxidants and other components such as pharmaceutically or biologically active substances, conferring desired properties to the finished chewing gum product.

30

Suitable bulk sweeteners include both sugar and non-sugar sweetening components. Bulk sweeteners typically constitute from about 5 to about 95% by weight of the

chewing gum, more typically about 20 to about 80% by weight such as 30 to 60% by weight of the gum.

Useful sugar sweeteners are saccharide-containing components commonly known in the chewing gum art including, but not limited to, sucrose, dextrose, maltose, dextrins, trehalose, D-tagatose, dried invert sugar, fructose, levulose, galactose, corn syrup solids, and the like, alone or in combination.

Sorbitol can be used as a non-sugar sweetener. Other useful non-sugar sweeteners include, but are not limited to, other sugar alcohols such as mannitol, xylitol, hydrogenated starch hydrolysates, maltitol, isomaltol, erythritol, lactitol and the like, alone or in combination.

High intensity artificial sweetening agents can also be used alone or in combination with the above sweeteners. Preferred high intensity sweeteners include, but are not limited to sucralose, aspartame, salts of acesulfame, alitame, saccharin and its salts, cyclamic acid and its salts, glycyrrhizin, dihydrochalcones, thaumatin, monellin, sterioside and the like, alone or in combination. In order to provide longer lasting sweetness and flavour perception, it may be desirable to encapsulate or otherwise control the release of at least a portion of the artificial sweetener. Techniques such as wet granulation, wax granulation, spray drying, spray chilling, fluid bed coating, coascervation, encapsulation in yeast cells and fibre extrusion may be used to achieve desired release characteristics. Encapsulation of sweetening agents can also be provided using another chewing gum component such as a resinous compound.

Usage level of the artificial sweetener will vary considerably and will depend on factors such as potency of the sweetener, rate of release, desired sweetness of the product, level and type of flavour used and cost considerations. Thus, the active level of artificial sweetener may vary from about 0.02 to about 8% by weight. When carriers used for encapsulation are included, the usage level of the encapsulated sweetener will be proportionately higher. Combinations of sugar and/or non-sugar sweeteners can be used in the chewing gum formulation processed in accordance

with the invention. Additionally, the softener may also provide additional sweetness such as with aqueous sugar or alditol solutions.

If a low calorie gum is desired, a low caloric bulking agent can be used. Examples of
5 low caloric bulking agents include polydextrose, Raftilose, Raftilin, fructooligosaccharides (NutraFlora[®]), palatinose oligosaccharides; guar gum hydrolysates (e.g. Sun Fiber[®]) or indigestible dextrins (e.g. Fibersol[®]). However, other low calorie-bulking agent can be used.

10 Further chewing gum additives which may be included in the chewing gum according to the present invention include surfactants and/or solubilisers, especially when pharmaceutically or biologically active ingredients are present. As examples of types of surfactants to be used as solubilisers in a chewing gum composition according to the invention reference is made to H.P. Fiedler, Lexikon der Hilfstoffe
15 für Pharmacie, Kosmetik und Angrenzende Gebiete, page 63-64 (1981) and the lists of approved food emulsifiers of the individual countries. Anionic, cationic, amphoteric or non-ionic solubilisers can be used. Suitable solubilisers include lecithin, polyoxyethylene stearate, polyoxyethylene sorbitan fatty acid esters, fatty acid salts, mono and diacetyl tartaric acid esters of mono and diglycerides of edible
20 fatty acids, citric acid esters of mono and diglycerides of edible fatty acids, saccharose esters of fatty acids, polyglycerol esters of fatty acids, polyglycerol esters of interesterified castor oil acid (E476), sodium stearyl sulfate, sodium lauryl sulfate and sorbitan esters of fatty acids and polyoxyethylated hydrogenated castor oil (e.g. the product sold under the trade name CREMOPHOR), block copolymers of
25 ethylene oxide and propylene oxide (e.g. products sold under trade names PLURONIC and POLOXAMER), polyoxyethylene fatty alcohol ethers, polyoxyethylene sorbitan fatty acid esters, sorbitan esters of fatty acids and polyoxyethylene stearic acid esters.

30 Particularly suitable solubilisers are polyoxyethylene stearates, such as for instance polyoxyethylene(8)stearate and polyoxyethylene(40)stearate, the polyoxyethylene sorbitan fatty acid esters sold under the trade name TWEEN, for instance TWEEN

20 (monolaurate), TWEEN 80 (monooleate), TWEEN 40 (monopalmitate), TWEEN 60 (monostearate) or TWEEN 65 (tristearate), mono and diacetyl tartaric acid esters of mono and diglycerides of edible fatty acids, citric acid esters of mono and diglycerides of edible fatty acids, sodium stearoyllatylate, sodium laurylsulfate, 5 polyoxyethylated hydrogenated castor oil, blockcopolymers of ethylene oxide and propyleneoxide and polyoxyethylene fatty alcohol ether. The solubiliser may either be a single compound or a combination of several compounds. In the presence of an active ingredient the chewing gum may preferably also comprise a carrier known in the art.

10

The chewing gum according to the present invention may contain aroma agents and flavouring agents including natural and synthetic flavourings e.g. in the form of natural vegetable components, essential oils, essences, extracts, powders, including acids and other substances capable of affecting the taste profile. Examples of liquid 15 and powdered flavourings include coconut, coffee, chocolate, vanilla, grape fruit, orange, lime, menthol, liquorice, caramel aroma, honey aroma, peanut, walnut, cashew, hazelnut, almonds, pineapple, strawberry, raspberry, tropical fruits, cherries, cinnamon, peppermint, wintergreen, spearmint, eucalyptus, and mint, fruit essence such as from apple, pear, peach, strawberry, apricot, raspberry, cherry, pineapple, 20 and plum essence. The essential oils include peppermint, spearmint, menthol, eucalyptus, clove oil, bay oil, anise, thyme, cedar leaf oil, nutmeg, and oils of the fruits mentioned above.

The chewing gum flavour may be a natural flavouring agent which is freeze-dried, 25 preferably in the form of a powder, slices or pieces of combinations thereof. The particle size may be less than 3 mm, such as less than 2 mm, more preferred less than 1 mm, calculated as the longest dimension of the particle. The natural flavouring agent may in a form where the particle size is from about 3 μm to 2 mm, such as from 4 μm to 1 mm. Preferred natural flavouring agents include seeds from a fruit 30 e.g. from strawberry, blackberry and raspberry.

Various synthetic flavours, such as mixed fruit flavours may also be used in the present chewing gum centres. As indicated above, the aroma agent may be used in quantities smaller than those conventionally used. The aroma agents and/or flavours may be used in an amount of from 0.01 to about 30% by weight of the final product depending on the desired intensity of the aroma and/or flavour used. Preferably, the content of aroma/flavour is in the range of from 0.2 to 3% by weight of the total composition.

In one embodiment the chewing gum composition comprises a pharmaceutically or biologically active substance. Examples of such active substances, a comprehensive list of which is found e.g. in WO 00/25598, which is incorporated herein by reference, include drugs, dietary supplements, antiseptic agents, pH adjusting agents, anti-smoking agents and substances for the care or treatment of the oral cavity and the teeth such as hydrogen peroxide and compounds capable of releasing urea during chewing. Examples of active substances in the form of agents adjusting the pH in the oral cavity include: acids, such as adipinic acid, succinic acid, fumaric acid, or salts thereof or salts of citric acid, tartaric acid, malic acid, acetic acid, lactic acid, phosphoric acid and glutaric acid and acceptable bases, such as carbonates, hydrogen carbonates, phosphates, sulphates or oxides of sodium, potassium, ammonium, magnesium or calcium, especially magnesium and calcium.

The gum centre of a coated chewing gum according to the invention can have any form, shape or dimension that permits the chewing gum centre to be coated using any conventional coating process. Accordingly, the gum centre may be e.g. in a form selected from a pellet, a cushion-shaped pellet, a stick, a tablet, a chunk, a pastille, a pill, a ball and a sphere.

The invention will now be described in further details in the following, non-limiting examples and figures wherein

Fig. 1 shows G^* and $\tan(d)$ versus frequency for synthesised polyisobutylene substitutes (PIB sub.1, PIB sub. 2, PIB sub. 3 and PIB sub. 4) including the two standards of PIB 1 and PIB 2,

- 5 Fig. 2 shows G' vs. osc. torque (micro N.m) for gum bases shown in Table 4 plus additional 2 conventional gum bases,

Fig. 3 shows $\tan(d)$ vs. osc. torque (micro N.m) for gum bases shown in Table 4 plus additional 2 conventional gum bases, and

10

Fig. 4 shows G' vs. osc. torque (micro N.m) for synthesised polyisobutylene substitutes and mixture hereof.

15 **EXAMPLE 1**

Evaluation of presently applied butyl rubber in chewing gum base

The elastomer portion of chewing gum base in a standard gum base typically comprises approximately 3-30% of the total material, and often consists of two polyisobutylene (PIB) fractions differing in molecular weight. A sample of PIB presently applied as elastomer in gum base, was analyzed by size exclusion chromatography (SEC) (see Table 1). The low molecular weight component of the PIB consisted of a material with a weight average molecular weight, M_w , of about 60,000 g/mol and a polydispersity (PDI) that varies in the range of 1.5-2.2.

25

TABLE 1: SEC Molecular weight data of currently applied PIB elastomers

Sample	M_n	M_w	PDI (M_w/M_n)
PIB 1	27,000	58,400	2.16
PIB 2	39,800	59,200	1.49

EXAMPLE 2

Preparation of polyisobutylene substitutes

- Poly(ϵ -caprolactone-co- δ -valerolactone) (denoted poly(CAP-co-VAL)) was prepared with a feed ratio of 60 mol % ϵ -caprolactone and 40 % δ -valerolactone (60 CAP:40 VAL). Poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) (denoted poly(CAP-co-VAL-TMC)) was prepared with a feed a ratio of 50 mol % ϵ -caprolactone, 40 mol % δ -valerolactone and 10 mol % trimethylene carbonate.
- The samples indicated in the below Table 2 were prepared for evaluation as polyisobutylene (PIB) substitutes.

TABLE 2

Sample	Composition	T _g (°C)	T _m (°C)	M _n (g/mol)	PDI
2169-37 PIB sub. 1	Poly(CAP-co-VAL))	-65	15	60,390	1.47
52-1 PIB sub. 2	Poly(CAP-co-VAL-TMC)	-65	10	51,190	1.63
A PIB sub. 3	Poly(CAP-co-VAL-TMC)	-60	16	50,780	1.44
B PIB sub. 4	Poly(CAP-co-VAL-TMC)	-60	16	53,340	1.56

Sample 2169-37 (PIB sub. 1) was further purified, and the M_n was subsequently measured to 54,850 g/mol indicating that the sample had started to degrade.

The synthesised samples were characterised as follows:

Characterisation

The structural characterisation of the above polymers was performed by routine ^{13}C and ^1H NMR spectroscopy. Spectra were acquired on a Bruker AC-300 (300 MHz) spectrometer using 5 mm O.D. tubes and deuterated chloroform as solvent with
5 internal standard tetramethylsilane (TMS). Sample concentrations were ~20% (w/v) for ^{13}C NMR and ~5% (w/v) for ^1H NMR spectra.

^{13}C NMR of poly(CAP-co-VAL) indicated that the feed ratio (60 mol% CAP and 40 mol % VAL) and the synthesised copolymer composition was approximately equal.

10

^{13}C NMR of the terpolymer poly(CAP-co-VAL-TMC) revealed a random structure and that the synthesised terpolymer composition and the feed ratio of monomers were approximately equal.

15 Size exclusion chromatography (SEC) experiments were performed to determine the molecular weights and polydispersities (PDI) of the polymeric materials. The SEC system is equipped with a Waters Alliance 2690 Separations Module, an on-line multiangle laser light scattering (MALLS) detector (MiniDAWN™, Wyatt Technology Inc.), an interferometric refractometer (Optilab DSP™, Wyatt
20 Technology Inc.) and one of two sets of PLgel™ (Polymer Laboratories Inc.) SEC columns. Each of the sets, consisting of two 3 μm or two 5 μm PLgel™ columns. The results are shown in Table 2.

Differential Scanning Calorimetry (DSC) was used to characterize the thermal
25 properties of the obtained biodegradable materials. The glass transition temperature (T_g) and melting temperature (T_m) were measured using either a Mettler DSC 30 or Perkin Elmer DSC-7. The samples were heated from -100°C to 100°C at a heating rate of 10°C/min. quenched, and heated again from -100°C to 100°C at the same rate. The results are shown in Table 2.

30

Rheological measurement were applied in order to select the most appropriate samples for up-scaling purpose.

In Fig. 1 G^* and $\tan(\delta)$ versus frequency are shown. These parameters are essential regarding texture properties of the final chewing gum. G^* is indicating the compactness/hardness of the chewing gum and $\tan(\delta)$, defining the ratio between loss modulus and storage modulus. The rheological evaluations were made using a rheometer, type AR1000 from TA instrument. The oscillation measurement is performed at a stress within the linear viscoelastic region and a temperature of 70 °C with a parallel plate system ($d=2,0$ cm, hatched).

PIB sub. 1 and PIB sub. 2 were chosen to be the best match and scaled up for further investigation in gum base and chewing gum.

The characteristics of the up-scaled samples PIB sub. 1 and PIB sub. 2 is shown in the below Table 3

TABLE 3

Sample	Composition	Tg (°C)	Tm (°C)	Mn (g/mol)	PDI
52-19 PIB sub. 1	poly(CAP-co-VAL)	-65	17	63,957	1.42
52-16T PIB sub. 2	poly(CAP-co-VAL-TMC)	-65	8	72,409	1.67

EXAMPLE 3

Preparation of polyisobutylene substitutes by means of mixing biodegradable polymers based on ϵ -caprolactone, δ -valerolactone and/or trimethylene carbonate.

This example demonstrates the possibility of creating biodegradable polymer substitutes for polyisobutylene (PIB) by means of mixing different molecular weight poly(ϵ -caprolactone-co- δ -valerolactone) and poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate).

Fig. 4 shows how a poly(ϵ -caprolactone-co- δ -valerolactone) with a molecular weight (Mn) of 18180 g/mol and a poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) with a molecular weight (Mn) of 76950 g/mol in a 50/50% mixture gives a rheological match to the standard PIB's.

The rheological evaluations were made using a rheometer, type AR1000 from TA Instrument. The oscillation measurement is performed at a frequency of 1 Hz and a temperature of 70 °C with a parallel plate system (d=2,0 cm, hatched).

EXAMPLE 4

Replacement of polyisobutylen in gum base with synthesised poly(ϵ -caprolactone-co- δ -valerolactone) and poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate)

The following experiment was performed in order to test synthesised poly(ϵ -caprolactone-co- δ -valerolactone) and poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) in a chewing gum base as replacements for polyisobutylene (PIB).

Thus, the synthesised poly(ϵ -caprolactone-co- δ -valerolactone) with Mn: 63,957 g/mol, Tg=-65 °C and Tm=17 °C (PIB sub. 1) and poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) with Mn: 72,409 g/mol, Tg=-65 °C and Tm=8 °C (PIB sub. 2) were tested in different gum base formulations.

The different gum base formulations were prepared in accordance with Table 4. The amounts in the compositions are given in percentage by weight. Samples 118 and 119 were prepared without heating during the mixing process except in the end of the mixing process where heat was applied in order to melt the softening system.

TABLE 4

	Standard (115)	PIB sub. 1 (116)	PIB sub. 2 (117)	PIB sub. 2 (118)*	PIB sub. 2 (119)*
Butyl	5%	5%	5%	5%	5%
Elastomer plasticizer	40%	40%	40%	40%	40%
Filler	16.5%	16.5%	16.5%	16.5%	16.5%
PIB sub. 1	-	14%	-	14%	-
PIB sub. 2	-	-	14%	-	14%
PIB low Mw	14%	-	-	-	-
softening system	24.5%%	24.5%	24.5%	24.5%	24.5%
Chewing gum no.	141	142	143	144	145

PIB sub. 1: poly(ϵ -caprolactone-co- δ -valerolactone), Mn=63,957 g/mol, Tg=-65°C, Tm=17°C.

PIB sub. 2: poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate), Mn=72,409 g/mol, Tg=-65°C, Tm=8°C,*Low temperature mix

5

The above gum bases were evaluated by means of rheology measurements; G' and tan (δ) vs. osc. torque (micro N.m) giving the linear viscoelastic region and thereby indicating the stability of the gum base structure. The results of these measurements are shown in Fig 2 and Fig. 3. As can be seen from the figures all the gum bases are very close to the standard gum base 115 and the two conventional gum bases included in the test set-up. The deviations are within the region of which gum bases can be described as having an acceptable quality.

The rheological evaluations were made using a rheometer, type AR1000 from TA Instrument. The oscillation measurement is performed at a stress within the linear viscoelastic region and a temperature of 70 °C with a parallel plate system (d=2,0 cm, hatched).

15

EXAMPLE 5

PIB replacement in a standard chewing gum formulation

The following experiment was conducted in order to test gum bases wherein polyisobutylene (PIB) was replaced with PIB sub. 1 and PIB sub. 2 in a standard peppermint chewing gum formulation. The standard peppermint chewing gum formulation was prepared in accordance with the below Table 5.

TABLE 5

	%
Gum base	38
Sorbitol powder	46
Maltitolsyrup	4
Xylitol powder	6
Peppermint oil	2.0
Carbamid	3.5
Peppermint powder	0.20
Aspartame	0.20
Acesulfame	0.10

Hardness was measured on the chewing gum samples indicated in Table 6. The hardness of the test samples were tested by an compression load test using an Instron instrument with a 4 mm DIA CYLINDER STAINLESS at a speed of 25 mm/min using a test distance of 3.5 mm into the chewing gum body.

TABLE 6

Chewing gum no.	Hardness (N); mean of 5
141 (14% PIB)	5.3
142 (14% PIB sub. 1)	5.1
143 (14% PIB. Sub. 2)	7.9
144 (14% PIB. Sub. 1)	5.9
145 (14% PIB. Sub. 2)	5.9

As can be seen from Table 6, the samples comprising either PIB sub. 1 or PIB sub. 2 are very close to the standard chewing gum (141) comprising 14% PIB. Hardness indicates that the initial chew is very close to the standard.

5

EXAMPLE 6

Sensorial evaluations

10 Test samples were evaluated by serving them to 10 trained panellists:

The following descriptive parameters were found when compared to the standard chewing gum (141).

15 Chewing gum no. 142: More cricky, waxy initial chew, but otherwise very close to standard 141.

Chewing gum no. 143: Harder/tough initial chew corresponds very well with hardness measurements, tacky, more cricky, otherwise very close to standard 141.

20

The samples mixed at lower temperatures seems to have better product quality regarding texture and tackiness (samples 144 and 145).

CLAIMS

1. An elastomer for chewing gum comprising a polyester polymer obtainable by the polymerisation of two or more different cyclic ester monomers, wherein the cyclic ester monomers have a low glass transition temperature (T_g) and the polyester polymer has a glass transition temperature (T_g) in the range from (-20°C) to (-80°C).
5
2. An elastomer for chewing gum of claim 1, wherein the cyclic ester monomers are selected from the group consisting of a 4-membered lactone, a 5-membered lactone, a 6-membered lactone, a 7-membered lactone, a 8-membered lactone, a 5-membered cyclic carbonate and a 6-membered cyclic carbonate.
10
3. An elastomer for chewing gum of claim 2, wherein the lactone is selected from the group consisting of β-propiolactone, γ-butyrolactone, δ-valerolactone, ε-caprolactone and 7-heptanolactone.
15
4. An elastomer for chewing gum of claim 2, wherein cyclic carbonate is an ethylene carbonate or a trimethylene carbonate.
- 20 5. An elastomer for chewing gum of claim 2, wherein the cyclic ester monomers are selected from the group consisting of ε-caprolactone, δ-valerolactone and trimethylene carbonate.
6. An elastomer for chewing gum of any of claims 1-5, wherein at least one cyclic ester monomer is ε-caprolactone.
25
7. An elastomer for chewing gum of any of claims 1-6, wherein the polyester polymer contains at least 50 mole% of ε-caprolactone.
- 30 8. An elastomer for chewing gum of any of claims 1-7, wherein the polyester polymer has a glass transition temperature (T_g) in the range from (-25°C) to (-75°C),

more preferably wherein the polyester polymer has a glass transition temperature (T_g) in the range from (-45°C) to (-75°C).

9. An elastomer for chewing gum comprising poly(ϵ -caprolactone-co- δ -valerolactone).

10. An elastomer for chewing gum according to claim 9 wherein the mol percentage of ϵ -caprolactone in said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 1 - 99 mol %.

11. An elastomer for chewing gum according to claim 10 wherein the mol percentage of ϵ -caprolactone in said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 40 - 80 mol %.

12. An elastomer for chewing gum according to claim 11 wherein the mol percentage of ϵ -caprolactone in said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 50 - 70 mol %.

13. An elastomer for chewing gum according to claim 12 wherein the mol percentage of ϵ -caprolactone in said poly(ϵ -caprolactone-co- δ -valerolactone) is about 60 mol %.

14. An elastomer for chewing gum according to claim 9 wherein the mol percentage of δ -valerolactone in said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 1 - 99 mol %.

15. An elastomer for chewing gum according to claim 14 wherein the mol percentage of δ -valerolactone in said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 20 - 60 mol %.

16. An elastomer for chewing gum according to claim 15 wherein the mol percentage of δ -valerolactone in said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 30 - 50 mol %.

17. An elastomer for chewing gum according to claim 16 wherein the mol percentage of δ -valerolactone in said poly(ϵ -caprolactone-co- δ -valerolactone) is about 40 mol %.
- 5 18. An elastomer for chewing gum according to claim 9 wherein the molecular weight (M_n) of said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 10,000 - 125,000 g/mol.
- 10 19. An elastomer for chewing gum according to claim 18 wherein the molecular weight (M_n) of said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 20,000 - 100,000 g/mol.
- 15 20. An elastomer for chewing gum according to claim 19 wherein the molecular weight (M_n) of said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 30,000 - 90,000 g/mol.
- 20 21. An elastomer for chewing gum according to claim 20 wherein the molecular weight (M_n) of said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of 40,000 - 80,000 g/mol.
22. An elastomer for chewing gum according to claim 9 wherein the glass transition temperature T_g of said poly(ϵ -caprolactone-co- δ -valerolactone) is less than 0°C.
- 25 23. An elastomer for chewing gum according to claim 22 wherein the glass transition temperature T_g of said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of (-40°C) - (-80°C).
- 30 24. An elastomer for chewing gum according to claim 23 wherein the glass transition temperature T_g of said poly(ϵ -caprolactone-co- δ -valerolactone) is in the range of (-50°C) - (-70°C).

25. An elastomer for chewing gum comprising poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate).
26. An elastomer for chewing gum according to claim 25 wherein the mol percentage
5 of ϵ -caprolactone in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 1 - 99 mol %.
27. An elastomer for chewing gum according to claim 26 wherein the mol percentage
10 of ϵ -caprolactone in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 20 - 80 mol %.
28. An elastomer for chewing gum according to claim 27 wherein the mol percentage
15 of ϵ -caprolactone in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 40 -60 mol %.
29. An elastomer for chewing gum according to claim 28 wherein the mol percentage
of ϵ -caprolactone in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is about 50 mol %.
- 20 30. An elastomer for chewing gum according to claim 25 wherein the mol percentage of δ -valerolactone in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 1 - 99 mol %.
31. An elastomer for chewing gum according to claim 30 wherein the mol percentage
25 of δ -valerolactone in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 20 - 60 mol %.
32. An elastomer for chewing gum according to claim 31 wherein the mol percentage
30 of δ -valerolactone in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 30 - 50 mol %.

33. An elastomer for chewing gum according to claim 32 wherein the mol percentage of δ -valerolactone in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is about 40 mol %.
- 5 34. An elastomer for chewing gum according to claim 25 wherein the mol percentage of trimethylene carbonate in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 1 - 50 mol %.
- 10 35. An elastomer for chewing gum according to claim 34 wherein the mol percentage of trimethylene carbonate in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 2 - 30 mol %.
- 15 36. An elastomer for chewing gum according to claim 25 wherein the mol percentage of trimethylene carbonate in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 5 - 15 mol %.
- 20 37. An elastomer for chewing gum according to claim 25 wherein the mol percentage of trimethylene carbonate in said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is about 10 mol %.
38. An elastomer for chewing gum according to claim 25 wherein the molecular weight (M_n) of said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 10,000 - 150,000 g/mol.
- 25 39. An elastomer for chewing gum according to claim 38 wherein the molecular weight (M_n) of said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 20,000 - 100,000 g/mol.
- 30 40. An elastomer for chewing gum according to claim 39 wherein the molecular weight (M_n) of said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 30,000 - 90,000 g/mol.

41. An elastomer for chewing gum according to claim 40 wherein the molecular weight (M_n) of said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of 40,000 - 80,000 g/mol.
- 5 42. An elastomer for chewing gum according to claim 25 wherein the glass transition temperature T_g of said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is less than 0°C.
- 10 43. An elastomer for chewing gum according to claim 42 wherein the glass transition temperature T_g of said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of (-40°C) - (-80°C).
- 15 44. An elastomer for chewing gum according to claim 43 wherein the glass transition temperature T_g of said poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) is in the range of (-50°C) - (-75°C).
- 20 45. An elastomer for chewing gum comprising a mixture of a poly(ϵ -caprolactone-co- δ -valerolactone) according to any of claims 9-24 and a poly(ϵ -caprolactone-co- δ -valerolactone-co-trimethylene carbonate) according to any of claims 25-44.
46. Gum base comprising elastomer and resin, wherein said elastomer comprises a biodegradable polymer.
- 25 47. Gum base according to claim 46, wherein said resin comprises a non-biodegradable resin.
48. Gum base according to claim 46 or 47, wherein said gum base comprises softener.
- 30 49. Gum base comprising elastomer and resin, wherein said elastomer comprises a biodegradable polymer according to any of the claims 1-45.

50. Gum base according to claim 49, wherein said resin comprises a non-biodegradable resin.

5 51. Gum base according to claim 49 or 50, wherein said gum base comprises softener.

52. Chewing gum comprising
elastomer and resin,
10 wherein said elastomer comprises a biodegradable polymer.

53. Chewing gum according to claim 52, wherein said resin comprises a non-biodegradable resin.

15 54. Chewing gum according to claim 52 or 53, wherein said gum base comprises softener.

55. Chewing gum comprising elastomer and resin, wherein said elastomer comprises a biodegradable polymer according to any of the claims 1-45.
20

56. Chewing gum according to claim 55, wherein said resin comprises a non-biodegradable resin.

57. Chewing gum according to claim 55 or 56, wherein said gum base comprises
25 softener.

58. Method of manufacturing a chewing gum, whereby at least one elastomer has been substituted by at least one biodegradable elastomer having rheologically properties substantially matching at least one conventional non-biodegradable
30 elastomer suitable for chewing gum.

59. Method according to claim 58, whereby the at least one biodegradable elastomer comprises a biodegradable polymer according to any of the claims 1-45.

1 / 4

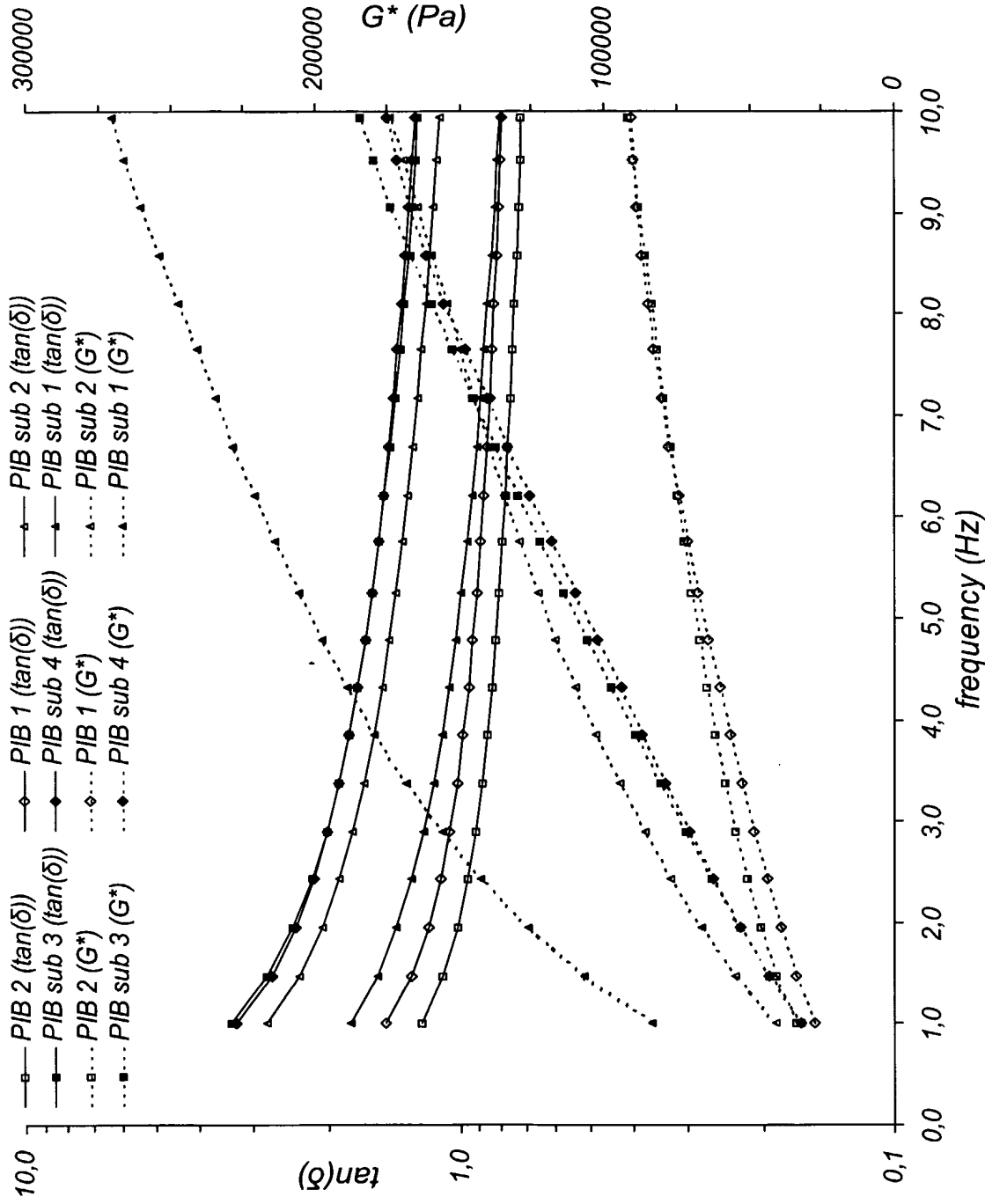


Fig. 1

2 / 4

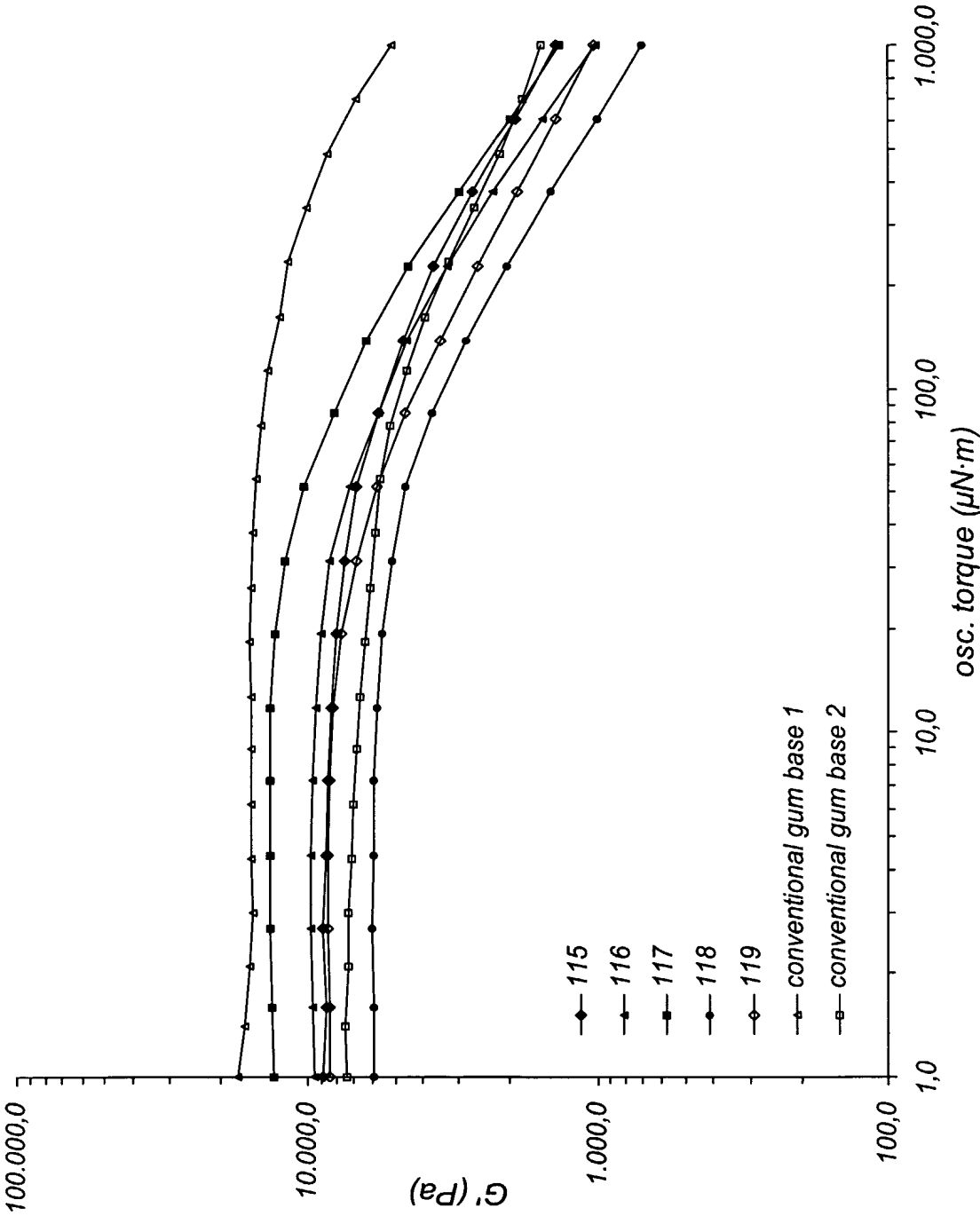


Fig. 2

3/4

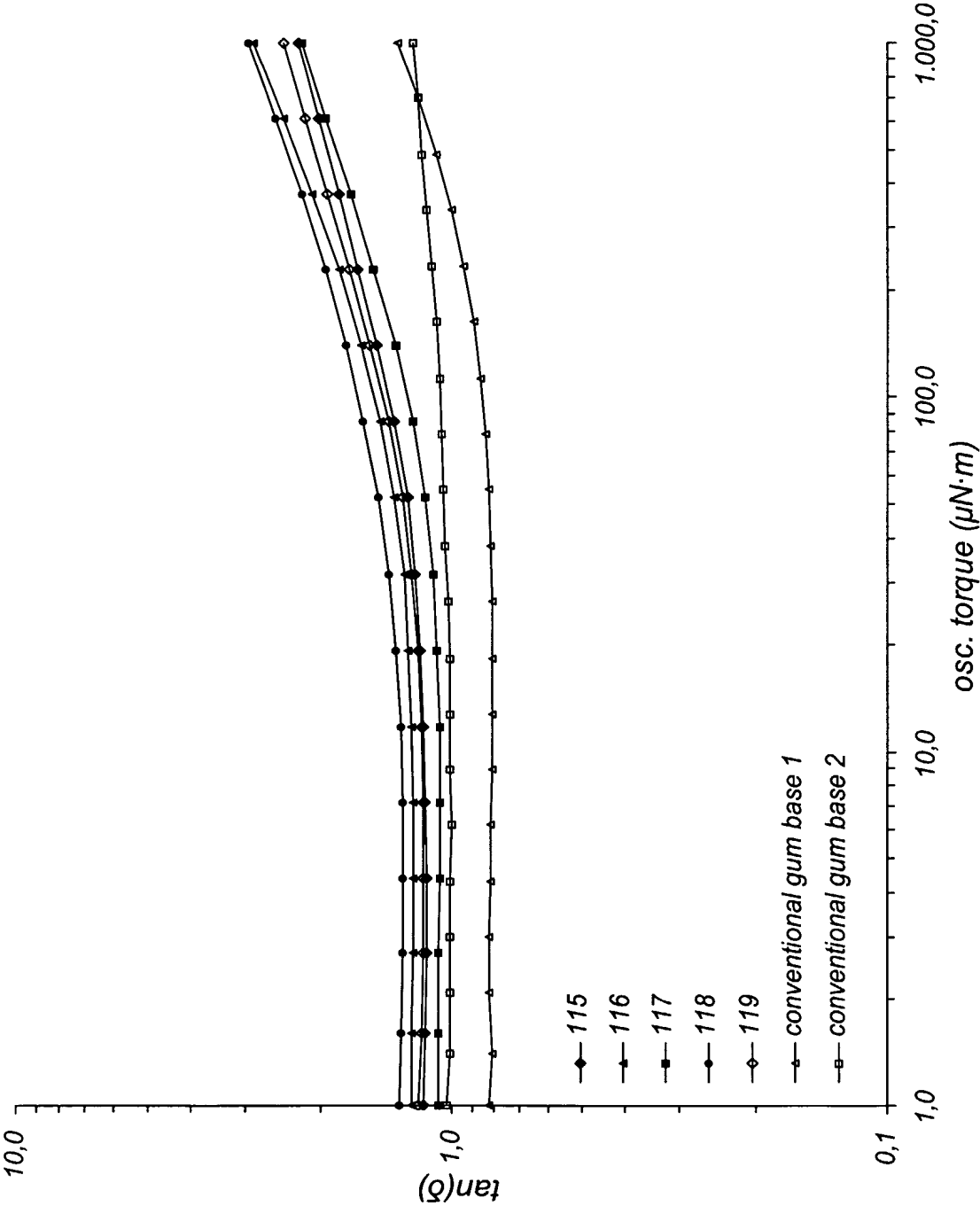


Fig. 3

4 / 4

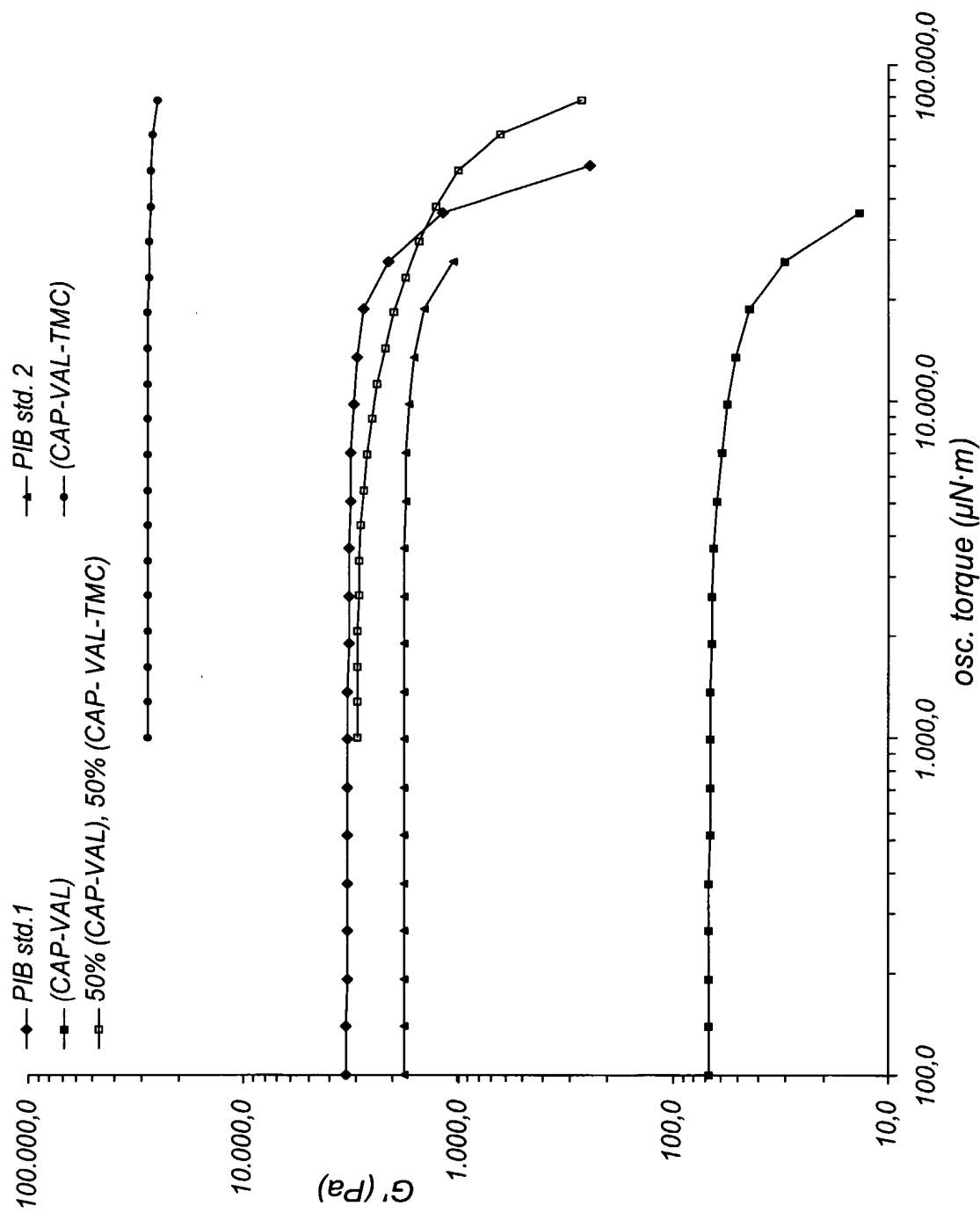


Fig. 4

INTERNATIONAL SEARCH REPORT

International Application No

PCT/DK 02/00203

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 7 A23G3/30 C08G63/08 C08G63/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A23G C08G

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, PAJ, EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 558 965 A (AMERICAN CYANAMID CO) 8 September 1993 (1993-09-08) column 9, line 16 - line 39; claims 1,2,4; example 2 column 4, line 34 - line 38 ---	1-9, 18, 22-25, 38, 42-44
X	US 5 672 367 A (GRIJPMAN DIRK WYBE ET AL) 30 September 1997 (1997-09-30) cited in the application column 1, line 29 - line 65; claims 1-8; examples 1,3 ---	1-6, 8, 9, 22-24, 46-59
X	US 5 530 074 A (JESSUP GEORGE ET AL) 25 June 1996 (1996-06-25) the whole document --- -/--	1-9, 18, 22-25, 38, 42-44



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

G document member of the same patent family

Date of the actual completion of the international search

11 July 2002

Date of mailing of the international search report

19/07/2002

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

Guyon, R

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/DK 02/00203

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 6 190 773 B1 (WATANABE YASUYUKI ET AL) 20 February 2001 (2001-02-20) the whole document ---	1-9
A	WO 94 11441 A (UNIV GRONINGEN ;GRIJPMA DIRK WYBE (NL); JOZIASSE CONRNELIS AARNOUD) 26 May 1994 (1994-05-26) the whole document ---	1-6
A	US 5 610 266 A (BUCHHOLZ BERTHOLD) 11 March 1997 (1997-03-11) column 2, line 33 - line 42; examples 1,2 ---	1
A	US 6 153 231 A (LI WEISHENG ET AL) 28 November 2000 (2000-11-28) the whole document ---	1
A	WO 00 19837 A (LI WEISHENG ;WRIGLEY W M JUN CO (US)) 13 April 2000 (2000-04-13) ---	
A	EP 0 500 098 A (HIMONT INC) 26 August 1992 (1992-08-26) the whole document ---	1
A,P	WO 01 47368 A (PATEL BHARAT KANAIYALAL ;GOLDBERG DANIEL (US); EATON ROBERT FRANCI) 5 July 2001 (2001-07-05) the whole document -----	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/DK 02/00203

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0558965	A	08-09-1993	US 5352515 A AT 195540 T AU 671976 B2 AU 3391793 A CA 2090565 A1 DE 69329210 D1 DE 69329210 T2 EP 0558965 A2 ES 2149784 T3 JP 6041310 A MX 9301047 A1 PL 297901 A1 US 5442016 A US 5530074 A US 5621050 A ZA 9301445 A	04-10-1994 15-09-2000 19-09-1996 09-09-1993 03-09-1993 21-09-2000 05-04-2001 08-09-1993 16-11-2000 15-02-1994 01-11-1993 15-11-1993 15-08-1995 25-06-1996 15-04-1997 23-09-1993
US 5672367	A	30-09-1997	NL 9401703 A EP 0711506 A2 FI 954867 A JP 8196214 A	01-05-1996 15-05-1996 15-04-1996 06-08-1996
US 5530074	A	25-06-1996	US 5442016 A US 5352515 A US 5621050 A AT 195540 T AU 671976 B2 AU 3391793 A CA 2090565 A1 DE 69329210 D1 DE 69329210 T2 EP 0558965 A2 ES 2149784 T3 JP 6041310 A MX 9301047 A1 PL 297901 A1 ZA 9301445 A	15-08-1995 04-10-1994 15-04-1997 15-09-2000 19-09-1996 09-09-1993 03-09-1993 21-09-2000 05-04-2001 08-09-1993 16-11-2000 15-02-1994 01-11-1993 15-11-1993 23-09-1993
US 6190773	B1	20-02-2001	JP 2000007789 A EP 0955331 A2	11-01-2000 10-11-1999
WO 9411441	A	26-05-1994	NL 9201949 A AU 5577894 A DE 69312312 D1 DE 69312312 T2 EP 0667885 A1 JP 8504850 T WO 9411441 A1	01-06-1994 08-06-1994 21-08-1997 08-01-1998 23-08-1995 28-05-1996 26-05-1994
US 5610266	A	11-03-1997	DE 3937272 A1 AT 159538 T DE 59010767 D1 DK 427185 T3 EP 0427185 A2 ES 2110403 T3 GR 3025838 T3 JP 3057374 B2 JP 3177423 A	16-05-1991 15-11-1997 27-11-1997 15-12-1997 15-05-1991 16-02-1998 30-04-1998 26-06-2000 01-08-1991

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/DK 02/00203

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 5610266	A		JP 11263830 A	28-09-1999
US 6153231	A	28-11-2000	WO 9858550 A1 AU 3502697 A GB 2341782 A ,B	30-12-1998 04-01-1999 29-03-2000
WO 0019837	A	13-04-2000	WO 0019837 A1 AU 9678298 A EP 1133240 A1	13-04-2000 26-04-2000 19-09-2001
EP 0500098	A	26-08-1992	IT 1249045 B CA 2061608 A1 DE 69216652 D1 DE 69216652 T2 DK 500098 T3 EP 0500098 A2 ES 2097225 T3 JP 5255488 A US 5359027 A	11-02-1995 22-08-1992 27-02-1997 12-06-1997 14-07-1997 26-08-1992 01-04-1997 05-10-1993 25-10-1994
WO 0147368	A	05-07-2001	AU 2608701 A WO 0147368 A1	09-07-2001 05-07-2001